

Peanut

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Scientific Name and Introduction: *Arachis hypogaea* L., the peanut or groundnut, is an annual herb of the Leguminosae family. Two subspecies are grown commercially, and four market types are of greatest economic importance. *A. hypogaea* subsp. *hypogaea* includes the market types runner and virginia, and *A. hypogaea* subsp. *fastigata* includes the market types spanish and valencia. In the U.S., in the 1980's, 70% of the peanuts grown were runners, while 20, 10 and < 1% were virginia, spanish and valencia market types, respectively (Knauff and Gorbett, 1989). According to the USDA national peanut tonnage report, U.S. production in 1999 included 78% runners, 18% virginia, 3% spanish and 3% valencia. The edible portion is a seed which develops underground inside a pod containing two to four seeds. Peanuts originated in South America and are now cultivated world wide with the majority of production in India, Asia and the U.S. In the U.S., about 60% of peanuts are utilized in a variety of food products, with the remainder used in approximately equal proportions for export and for production of edible oil. Major chemical constituents of peanuts are oil (44 to 56%) and protein (22 to 33%), with a large influence of environment, genotype and maturity on their concentrations (Holaday and Pearson, 1974).

Quality Characteristics and Criteria: Shelled peanuts should be properly sized to meet market type, be free of misshapen or under-developed kernels and be free of any shell or foreign material and off-odor or flavor. Raw peanuts should be surrounded by a tan, pink or red- colored seed coat (testa) that fully encapsulates the seed, and the interior color of each half-seed should be ivory. Moisture content for in-shell peanuts should be < 10% to prevent mold growth (Diener and Davis, 1977). Prior to shelling, peanuts should contain 7 to 10% moisture to reduce splitting and kernel breakage during milling. After milling, moisture content for maximum shelf-life is \leq 7%. Seed may be stored at ambient temperature for up to 11 years with good viability if seed moisture content is < 3.3% (Cheng et al., 1997). Peanuts marketed without seed coats (blanched) should have an ivory colored raw kernel. Peanuts are most commonly consumed following roasting that may be accomplished in-shell or after shelling. Roasted peanut kernels should be light-yellow in color, free of external oil, contain < 6% moisture and be free of dark-colored kernels. Peanut seed, and particularly peanut seed coats, are a source of resveratrol, a compound that reduces cardiovascular disease and cancer incidence. Resveratrol ranges from 0.02 to 1.79 $\mu\text{g/g}$, compared to 0.6 to 8.0 $\mu\text{g/ml}$ in red wines (Sanders et al., 2000). Fatty acid composition of peanut oil is predominantly oleic and linoleic acid, found in roughly equal amounts, and making up 80% of total fatty acids. Certain genotypes may contain substantially more oleic than linoleic acid, with ratios as high as 40:1 (oleic:linoleic). Peanuts with a high oleic:linoleic acid ratio are less susceptible to oxidative deterioration and off-flavor development caused by oxidative cleavage of polyunsaturated fatty acids. The ratio of oleic to linoleic is influenced primarily by genotype, but interactions exist between genotype and the environment.

Horticultural Maturity Indices: Peanuts are an indeterminate plant, with flowering followed by underground seed development occurring over a range of time. Assessment of peanut maturity should be conducted using multiple plants at various locations within a field. Peanut maturity may be judged by the shell-out procedure, involving separation of peanut seed into mature or immature categories. Using the shell-out procedure, a peanut is considered mature if the inner hull is brown and the seed coat is pink to red. Optimum maturity is reached for runner and spanish types when 75 to 80% of the inner hull has turned brown, and for virginia types when 65% of the seed coat has turned deep pink in color. Runner peanut maturity can be determined by a hull-scrape method, in which maturity profiles for samples are

estimated based on degree of change in pod mesocarp from white to brown to black (Williams and Drexler, 1981).

Grades, Sizes and Packaging: U.S. grade standards and industry grade standards from the American Peanut Shellers Association (ASPA) exist for shelled spanish, shelled runner, shelled virginia and in-shell virginia peanuts. A comparison of tolerances as provided by the American Peanut Shellers Association Official Trade Rules for shelled peanuts based primarily on size, peanuts of other types, amount of split or broken kernels, freedom from foreign material, damage, minor defects and in some cases moisture percent is presented in Tables 1 to 4. Tolerances for in-shell virginia peanuts based on maturity, freedom from loose shelled peanuts, discoloration of shell, dirt, shell and other foreign material and degree of kernel fill inside the shell are compared.

Table 1. Comparison of screen sizes and tolerances (by weight) for American Peanut Shellers Association and USDA Grades for shelled spanish peanuts.

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Table 3. Comparison of screen sizes and tolerances (by weight) for American Peanut Shellers Association and USDA Grades for shelled Virginia peanuts

| Grade & Minimum Screen Size | % Fall-Thru Prescribed Screens | % Other Types | % Sound Split or Broken Kernels | % Damage | % Damage & Minor Defects | % Foreign Material | % Moisture | Count Per Pound Maximum |
|--|--------------------------------|---------------|---------------------------------|----------|--------------------------|--------------------|------------|-------------------------|
| EXTRA LARGE 20/64 x 1" Slot APSA - US | 3.00 | 0.75 | 3.00 | 1.00 | 1.75 | 0.10 | ---- | **512 |
| MEDIUM VIRGINIA 18/64 x 1" Slot APSA - US | 3.00 | 1.00 | 3.00 | 1.25 | 2.00 | 0.10 | ---- | **640 |
| NO. 1 VIRGINIA 15/64 x 1" Slot APSA - US | 3.00 | 1.00 | 3.00 | 1.25 | 2.00 | 0.10 | ---- | **864 |
| VIRGINIA SPLITS 20/64" Round APSA - US | *3.00 | 2.00 | Not Less Than 90.00 | ---- | 2.00 | 0.20 | ---- | ---- |
| NO. 2 VIRGINIA 17/64" Round APSA - US | *6.00 | 2.00 | As Graded | ---- | 2.50 | 0.20 | ---- | ---- |
| * (APSA-US) - Includes both sound split and broken and sound whole kernels which pass through prescribed screens. | | | | | | | | |
| “With Splits” - Each of the above APSA whole grades may be certified as “with splits” providing all requirements of the grade are met: except a tolerance of 15% is allowed for split kernels of which not more than 3% will pass through 17/64" round screen. | | | | | | | | |

** U.O.S. - Unless Otherwise Specified

Table 4. Comparison of screen sizes and tolerances (by weight) for American Peanut Shellers Association and USDA Grades for in-shell Virginia peanuts.

| Grade & Minimum Screen Size | % Fall-Thru Prescribed Screen | % Cracked or Broken Shells, Pops, Paper & Foreign Material | % Damaged Kernels | Count Per Pound |
|-----------------------------------|-------------------------------|--|-------------------|----------------------------------|
| JUMBO 37/64 x 3" APSA US | 5.00 5.00 | *10.00 *10.00 | 3.50 3.50 | **176 (U.O.S.) **176 (U.O.S.) |
| FANCY 32/64 x 3" APSA US | 5.00 5.00 | *11.00 *11.00 | 4.50 4.50 | **225 (U.O.S.) **225 (U.O.S.) |

* (APSA-US) - Not more than .5% shall be allowed as dirt or other foreign material.

** U.O.S. - Unless Otherwise Specified.

Raw in-shell peanuts are typically stored as “farmer stock peanuts” in flat, ventilated warehouses or grain bins in bulk or, less commonly in 50 lb burlap bags, for a week to 10 mo prior to shelling (Smith, Blankenship and McIntosh, 1995). After shelling, raw peanuts are often shipped in bulk containers, but may be packaged in burlap or nylon tote bags of various sizes. Peanuts for human consumption must be free of visible *Aspergillus flavus* mold, contain less than 15 ppb aflatoxin. Lots imported into the U.S. may be designated “segregation 1” or “segregation 2” depending on degree of kernel damage and concealed damage from rancidity, mold or decay. Segregation 1 peanuts may not contain more than 2.00% damaged kernels nor more than 1.00% concealed damage while segregation 2 peanuts may contain more than 2.00% damaged or more than 1.00% concealed damage. Segregation 3 peanuts are those which contain visible *Aspergillus flavus* mold or 15 ppb or more aflatoxin.

Optimum Storage Conditions: In-shell farmers stock peanuts dried to about 7.5% moisture. If stored at 10 °C (50 °F), these can be stored for up to 10 mo without significant quality loss (Davidson et al., 1982). High losses in milling quality may occur if peanuts are dried to below 7% moisture or if kernel temperature is below 7 °C (44.6 °F) during shelling (McIntosh and Davidson, 1971). Peanut moisture contents > 10% should be avoided to prevent mold growth (Diener and Davis, 1977). Adequate ventilation in a warehouse storage facility, preferably providing one air change every 3 min, is also desirable to prevent excess moisture and heat from accumulating in the storage facility (Smith and Davidson, 1982). Quality of raw shelled peanuts can be maintained for at least 1 year at 1 to 5 °C (33.8 to 41 °F) with moisture contents < 7%, or for 2 to 10 years at -18 °C (0 °F) and < 6 % moisture. Maintaining RH between 55 and 70% at 1 to 5 °C (33.8 to 41 °C) will maintain peanut moisture content at 7 to 7.5%. Careful handling of peanuts equilibrated to < 5 °C (41 °F) is necessary to prevent bruising and subsequent oil seepage from damaged cells within the cotyledon. Upon removal of raw shelled peanuts from refrigerated or frozen storage, equilibration to ambient temperature should be gradual in conditioning rooms, with RH/temperature/air-flow adjusted to prevent moisture condensation onto peanuts.

Controlled Atmosphere Considerations: Low O₂ storage shows promise for delaying rancid flavor development and insect infestation (Slay et al., 1985). High CO₂ storage appears to limit growth of *Aspergillus flavus* in short duration storage of high moisture, non-cured peanuts. Peanuts at 20% moisture stored at 0.6 to 3 °C (33 to 37.4 °F) in a high CO₂ environment had acceptable quality for 4 days, but deteriorated after 8 days of storage (Moseley et al., 1971). For longer term storage of high moisture, shelled peanuts under ambient temperature conditions, O₂ < 1.5% were required to slow *Aspergillus flavus* growth, but no CA totally eliminated aflatoxin production (Wilson and Jay, 1976).

Retail Outlet Display Considerations: Peanuts are normally marketed at ambient temperature conditions. Use of low O₂ and preventing excessive exposure to light is recommended. In-shell peanuts may be displayed and marketed in bulk containers. Exposure to moisture or high RH should be avoided.

Chilling Sensitivity: Prior to or during harvest, and prior to postharvest drying, exposure to chilling temperatures of 0.9 to 1.6 °C (33.6 to 34.9 °F) caused increased ethanol production, and the effect was greater for small as compared to large seed within a genotype. This was accompanied by increased seed leachate, suggested by the authors to be indicative of induction of anaerobic respiration and cell membrane damage (Singleton and Pattee, 1989). Following postharvest drying and during storage peanuts do not appear to be sensitive to chilling temperatures and may be stored at or below freezing.

Ethylene Production and Sensitivity: Peanut seeds exhibit dormancy periods following harvest of 63 to 84 days, varying with genotype and temperature during pod maturation and storage. Soaking seeds in 50 to 200 µg GA₃ or ethephon per mL of solution was effective in breaking dormancy (Kapur et al., 1990). Non-dormant peanuts exhibit a climacteric-like rise in ethylene production during seed germination (Whitehead and Nelson, 1992).

Respiration Rates: Properly cured peanuts in storage exhibit a relatively low rate of respiration. During harvest and prior to curing, especially for high moisture peanuts, respiration rates may be substantial and significant losses in quality can ensue. Freshly harvested peanuts should be dried soon after harvest to < 10% moisture to assure optimum quality.

Physiological Disorders: Shriveled seed trait has been identified as a heritable condition for peanuts. Seed mature normally but appear shriveled, and thus appear to have been harvested while immature. Seeds of shriveled lines exhibit up to 67% less oil, double the amount of sucrose, and defatted meal contained less protein (Jakkula et al., 1997). Improper curing of peanuts results in loss of quality and off-flavor development. Freezing temperatures occurring during harvest while peanuts are still windrowed (Singleton and Pattee, 1991), or curing at too high a temperature (Sanders et al., 1989) resulted in fermented, fruity off-flavor. Effects of improper curing are greatest on smaller seed, perhaps indicating greater effect on immature seed (Sanders et al., 1990).

Postharvest Pathology: Peanuts are susceptible to infection by various molds and fungi, and a combination of storage at 1 to 5 °C (33.8 to 41 °F) and reduction of moisture content < 7.5% may be effective in reducing mold and fungi growth in storage. The presence of toxic fungal metabolites (mycotoxins) are a particular concern. The name aflatoxin refers to four metabolites found in contaminated peanuts and designated aflatoxin B₁, B₂, G₁ and G₂. Aflatoxins B₁ and B₂ are metabolites of *Aspergillus flavus* and all four aflatoxins may be produced by *Aspergillus parasiticus* (Cole et al., 1995). A fifth mycotoxin which is somewhat less toxic than aflatoxin and is produced by *Aspergillus flavus*, other *Aspergillus* species, and several species of *Penicillium*, is cyclopiazonic acid (Dorner et al., 1985). Pre-harvest conditions favoring aflatoxin contamination are high temperatures and drought stress during the last 3 to 6 weeks of the growing season (Cole et al., 1989). Late season irrigation may be effective in reducing aflatoxin contamination (Dorner et al., 1989). When aflatoxin contamination occurs, it is common for most of the harvested seed to be free of contamination with only a few highly contaminated seed. Although monitoring at the point of sale for *Aspergillus flavus* is mandated by the USDA Peanut Marketing Agreement, and detection of 15 ppb aflatoxin or more leads to positive aflatoxin identification, the irregular distribution of infection may lead to false negative designations. Storage conditions to deter growth of the causal organisms and subsequent metabolic production of the mycotoxins primarily involve prevention of rehydration during storage. Decontamination of contaminated lots is most effectively done with electronic color sorting, although size and density separation may also be effective in removal of the most susceptible underdeveloped seed.

Quarantine Issues: Importation of peanut seed into the U.S. for planting is prohibited from Burkino, Faso, the People's Republic of China, Cote de'Ivoire, India, Indonesia, Ivory Coast, Japan, the Phillipines, Senegal, Thailand and Taiwan because of peanut stripe virus. During import of peanuts, all lots must be labeled with a positive lot identification and must meet the requirements for segregation 1 peanuts if used for human consumption. Peanuts with visible *Aspergillus flavus* mold, or those containing ≥ 15 ppb aflatoxin, may not be used for edible purposes; they may be used for oil stock. Such peanuts may be blanched and re-separated into aflatoxin-negative lots which may be used for edible purposes.

Special Considerations: Peanuts are a major allergenic food among adults and children in the U.S. (Taylor, 1992). Allergen activity has been identified for at least six allergens by phage display technology, and are known to be in association with the two major storage proteins, arachin and conarachin, and in profilin (Kleber et al., 1999). Allergens may also be present in refined peanut oil (Olszewski et al., 1998). Due in large part to allergenicity, any food product containing peanuts or peanut oil must be labeled as such. Careful handling of cured farmers stock peanuts to prevent breakage of shells reduces the risk for spread of fungal contamination and maintains the grade. Conveyors, cleaners, sizers

and other handling equipment should be padded where appropriate and properly maintained to prevent excessive breakage during handling. Once shelled and roasted, peanuts should be handled carefully to prevent separation of the half kernels and breakage since splits and pieces are more susceptible to oxidative deterioration and rancidity development. Peanuts will absorb lipophilic volatiles from their surroundings, or from inappropriate packaging that can induce off-flavors. Absorption of ammonia can cause darkening of nutmeats.

References:

- Cheng, H.Y., G.H. Zheng, X.M. Jing, T.Y. Kuang, P.S. Tang, K.L. Tao, M.D. Zhou and N.X. Duan. 1997. Storage of peanut seeds with low moisture content for 11 years in ambient temperature. *Plant Gen. Resour. Newsletter* 110:35-40.
- Cole, R.J., T.H. Sanders, J.W. Dorner and P.D. Blankenship. 1989. Environmental conditions required to induce preharvest aflatoxin contamination of groundnuts: Summary of six years research. In: *Aflatoxin contamination of groundnut. Proc. Intl. Wkshp, ICRISAT Cntr, Patancheru, Peru*, pp. 279-287.
- Cole, R.J., J.W. Dorner and C.C. Holbrook. 1995. Advances in mycotoxin elimination and resistance. In: H.E. Pattee and H.T. Stalker (eds) *Adv. Peanut Sci., Amer. Peanut Res. Ed. Soc., Stillwater OK*, pp. 456-474.
- Davidson, J.I., T.B. Whitaker and J.W. Dickens. 1982. Grading, cleaning, storage, shelling and marketing of peanuts in the United States. In: H.E. Pattee and C.T. Young (eds) *Peanut Sci. Technol., Amer. Peanut Res. Ed. Soc., Yoakum TX*, pp. 571-623.
- Diener, U.L. and N.D. Davis. 1977. Aflatoxin formation in peanuts by *Aspergillus flavus*. *Auburn Univ. Agric. Exp. Stn. Bull.* 493.
- Dorner, J.W., R.J. Cole and L.G. Lomax. 1985. The toxicity of cyclopiazonic acid. In: J. Lacey (ed) *Trichothecenes and Other Mycotoxins*, John Wiley and Sons, NY, pp. 529-535.
- Dorner, J.W., R.J. Cole, T.H. Sanders and P.D. Blankenship. 1989. Interrelationship of kernel water activity, soil temperature, maturity and phytoalexin production in pre-harvest aflatoxin accumulation of drought-stressed peanuts. *Mycopathologia* 105:117-128.
- Holaday, C.E. and J.L. Pearson. 1974. Effects of genotype and production area on the fatty acid composition, total oil and total protein in peanuts. *J. Food Sci.* 39:1206-1209.
- Jakkula, L.R., S.F. O'Keefe, D.A. Knauff and K.J. Boote. 1997. Chemical characterization of a shriveled seed trait in peanut. *Crop Sci.* 37:1560-1567.
- Kapur, A., J. Kaur, H.L. Sharma and H. Singh. 1990. Preconditioning of peanut (*Arachis hypogaea*) seeds to release dormancy. *Ann. Biol. Ludhiana* 6:141-145.
- Kleber, J.T., R. Cramer, U. Appenzeller, M. Cshlaak and W.M. Becker. 1999. Selective cloning of peanut allergens, including profilin and 2S albumins, by phage display technology. *Int. Arch. Allergy Immun.* 119:265-274.
- Knauff, D.A. and D.W. Gorbet. 1989. Genetic diversity among peanut cultivars. *Crop Sci.* 29:1417-1422.
- McIntosh, F.P. and J.I. Davidson. 1971. Effect of temperature on shelling runner- and spanish-type peanuts. *USDA, ARS 52-65, U.S. Govt. Print. Office, Washington DC*.
- Moseley, Y.C., H.B. Manbeck, G.L. Barnes and G.L. Nelson. 1971. Controlled atmosphere for short duration storage of peanuts before drying. *Trans. Amer. Soc. Agr. Eng.* 14:206-210.
- Olszewski, A., L. Pons, F. Moutete, G.I. Aimone, G. Kanny, D.A. Moneret-Vautrin and J.L. Gueant. 1998. Isolation and characterization of proteic allergens in refined peanut oil. *Clin. Exper. Allergy* 28:850-859.
- Sanders, T.H., J.R. VerCELLOTTI, P.B. Blankenship, K.L. Crippen and G.V. Civile. 1989. Interaction of maturity and curing temperature on descriptive flavor of peanuts. *J. Food Sci.* 54:1066-1069.
- Sanders, T.H., P.B. Blankenship, J.R. VerCELLOTTI and K.L. Crippen. 1990. Interaction of curing temperature and inherent maturity distribution in descriptive flavor of commercial grade sizes of Florunner peanuts. *Peanut Sci.* 17:85-89.

- Sanders, T.H., R.W. McMichael and K.W. Hendrix. 2000. Occurrence of resveratrol in edible peanuts. *J. Agric. Food Chem.* 48:1243-1246.
- Singleton, J.A. and H.E. Pattee. 1989. Effect of chilling injury on windrowed peanuts. *Peanut Sci.* 16:51-54.
- Singleton, J.A. and H.E. Pattee. 1991. Peanut moisture/size, relation to freeze damage and effect of drying temperature on volatiles. *J Food Sci.* 56:579-581.
- Slay, W.O., W.G. Ferguson and J.A. Pomplin. 1985. Some effects of conventional and low-oxygen atmosphere storage and processing methods on Florunner peanut seed. *Peanut Sci.* 12:8-11.
- Smith, J.S., P.D. Blankenship and F.P. McIntosh. 1995. Advances in peanut handling, shelling and storage from farmer stock to processing. In: H.E. Pattee and H.T. Stalker (eds) *Adv. Peanut Sci., Amer. Peanut Res. Edu. Soc., Stillwater OK*, pp. 500-527.
- Smith, J.S. and J.I. Davidson. 1982. Psychometrics and kernel moisture content as related to peanut storage. *Trans. Amer. Soc. Agric. Eng.* 25:231-236.
- Taylor, S.L. 1992. Chemistry and determination of food allergens. *Food Tech.* 46:146-152.
- Whitehead, C.S. and R.M. Nelson. 1992. Ethylene sensitivity in germinating peanut seeds: The effect of short-chain fatty acids. *J. Plant Physiol.* 139:479-483.
- Williams, E.J. and J.S. Drexler. 1981. A non-destructive method for determining peanut pod maturity. *Peanut Sci.* 8:134-141.
- Wilson, D.A. and E. Jay. 1976. Effect of controlled atmosphere storage on aflatoxin production in high moisture peanuts (groundnuts). *J. Stored Prod. Res.* 12:97-100.